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U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER 3ER

**TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371**

332013-024

U.S. APPLICATION NO. (If known, see 37 C.F.R. 1.5)

09/508378

INTERNATIONAL APPLICATION NO.
PCT/FR99/01680INTERNATIONAL FILING DATE
09 July 1999PRIORITY DATE CLAIMED
10 July 1998

TITLE OF INVENTION
SOLID POLYMER ELECTROLYTE AND PREPARATION METHODS

APPLICANT(S) FOR DO/EO/US
Jean-Yves SANCHEZ, Fannie ALLOIN

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This is an express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and the PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US)
6. ☒ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
- ☐ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☐ have not been made and will not be made.
- ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☐ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern other document(s) or information included:

11. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A FIRST preliminary amendment.
☐ A SECOND or SUBSEQUENT preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information:
International Search Report (in French)

U.S. APPLICATION NO. (If known, see 37 CFR 1.501) 097508378		INTERNATIONAL APPLICATION NO PCT/FR99/01680		ATTORNEY'S DOCKET NUMBER 032013-024	
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17. <input checked="" type="checkbox"/> The following fees are submitted:				CALCULATIONS	PTO USE ONLY																
Basic National Fee (37 CFR 1.492(a)(1)-(5)): Search Report has been prepared by the EPO or JPO \$840.00 (970) International preliminary examination fee paid to USPTO (37 CFR 1.482) \$670.00 (956) No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2)) \$690.00 (958) Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$970.00 (960) International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4) \$96.00 (962)																					
ENTER APPROPRIATE BASIC FEE AMOUNT =				\$ 840.00																	
Surcharge of \$130.00 (154) for furnishing the oath or declaration later than months from the earliest claimed priority date (37 CFR 1.492(e)).				\$																	
<table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <th style="width: 20%;">Claims</th> <th style="width: 20%;">Number Filed</th> <th style="width: 20%;">Number Extra</th> <th style="width: 20%;">Rate</th> </tr> <tr> <td>Total Claims</td> <td>36 -20 =</td> <td>16</td> <td>X\$18.00 (966)</td> </tr> <tr> <td>Independent Claims</td> <td>5 -3 =</td> <td>2</td> <td>X\$78.00 (964)</td> </tr> <tr> <td colspan="3">Multiple dependent claim(s) (if applicable)</td> <td>+ \$260.00 (968)</td> </tr> </table>				Claims	Number Filed	Number Extra	Rate	Total Claims	36 -20 =	16	X\$18.00 (966)	Independent Claims	5 -3 =	2	X\$78.00 (964)	Multiple dependent claim(s) (if applicable)			+ \$260.00 (968)	\$ 288.00	
Claims	Number Filed	Number Extra	Rate																		
Total Claims	36 -20 =	16	X\$18.00 (966)																		
Independent Claims	5 -3 =	2	X\$78.00 (964)																		
Multiple dependent claim(s) (if applicable)			+ \$260.00 (968)																		
				\$ 156.00																	
				\$ 0.00																	
TOTAL OF ABOVE CALCULATIONS =				\$ 1284.00																	
Reduction for 1/2 for filing by small entity, if applicable. Verified Small Entity statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28).				\$ 0.00																	
SUBTOTAL =				\$ 1284.00																	
Processing fee of \$130.00 (156) for furnishing the English translation later than months from the earliest claimed priority date (37 CFR 1.492(f)).				\$ 0.00																	
TOTAL NATIONAL FEE =				\$ 1284.00																	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 (581) per property +				\$ 0.00																	
TOTAL FEES ENCLOSED =				\$ 1284.00																	
				Amount to be: refunded	\$																
				charged	\$																

a. ☒ A check in the amount of \$ 1,284.00 to cover the above fees is enclosed.


b. ☐ Please charge my Deposit Account No. 02-4800 in the amount of \$_____ to cover the above fees. A duplicate copy of this sheet is enclosed.

c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 02-4800. A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

E. Joseph Gess
 BURNS, DOANE, SWECKER & MATHIS, L.L.P.
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 SIGNATURE
 E. Joseph Gess
 NAME
28,510
 REGISTRATION NUMBER

Patent
Attorney's Docket No. 032013-024

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application of)
)
Jean-Yves SANCHEZ et al.) Group Art Unit: Unassigned
)
Application No.: Unassigned) Examiner: Unassigned
)
Filed: March 10, 2000)
)
For: SOLID POLYMER ELECTROLYTE)
AND PROCESSES FOR ITS)
PREPARATION)

PRELIMINARY AMENDMENT

Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

Prior to an examination on the merits, please first amend the above-identified
application as follows:

IN THE SPECIFICATION

Page 1, after the title, please insert

--BACKGROUND OF THE INVENTION

Field of the Invention--.

Page 1, after line 11, please insert

--Description of the Related Art--.

Page 2, line 6, please delete "lithium/ion" and insert therefor --lithium ion--.

Page 3, after line 31, please insert

--SUMMARY OF THE INVENTION--.

Page 4, after line 23, please insert

--DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS--.

Page 7, line 17, please delete "kelvins" and insert therefor --Kelvin--; and

line 18, insert --temperature-- after "transition".

Page 8, line 13, delete "hexamethyl" and insert therefor --hexamethylene--.

Page 9, line 29, delete "(CH₂)_n- NH-CO-CH=CH₂" and insert therefor

--(CH₂)_n-NH-CO-CH=CH₂--;

line 31, please delete "alkyl" and insert therefor --alkylene--; and

line 35, please delete "methoxy" and insert therefor --methylether-- .

Page 10, line 15, please delete "methoxy" and insert --methylether--.

Page 14, line 24, delete "(C_nF_{2n}SO₃Li)" and insert therefor --(C_nF_{2n}SO₃Li)--.

Page 18, line 22, delete "(CP/EC/LiC₁₀)" and insert therefor --(PC/EC/LiC₁₀)--.

Page 20, line 3, please delete "making" and insert therefor --make--; and
line 5, delete "MAN" and insert therefor --MANHP--.

Page 21, line 6, please delete "insoluble" and insert therefor --insolubles--;
line 25, delete "angstroms" and insert therefor --Angströms--;
line 26, delete "Mass molar" and insert therefor --Molar--; and
line 30, delete "hexamethyl" and insert therefor --hexamethylene--.

Page 22, line 6, please delete "insoluble" and insert therefor --insolubles--; and
line 32, delete "methoxy" and insert therefor --methylether--.

Page 24, line 4, please delete "group" and insert therefor --groups--;
line 5, delete "insoluble" and insert therefor --insolubles--;
line 28, delete "group" and insert therefor --groups--; and
line 37, delete "Man" and insert therefor --MAN--.

Page 25, line 6, please delete "group" and insert therefor --groups--; and
line 7, delete "insoluble" and insert therefor --insolubles--.

Page 26, table, last row, please delete "PEG7400" and insert therefor --PEG8000--.

Page 29, line 20, please delete "Li/Li+" and insert therefor --Li/Li⁺-- and

line 22, delete "Li/Li+" and insert therefor --Li/Li⁺--.

Page 31, line 28, please delete "CA" and insert therefor --HA--.

Page 35, line 19, please delete "Example 19" and insert therefor --Example 18--.

IN THE CLAIMS

Kindly amend the following claims:

Claim 2, third from last line, please delete "of the" and insert therefor --of a--.

Claim 4, lines 1 and 2, please delete "either of Claims 1 and 2" and insert therefor
--Claim 1--.

Claim 8, line 3, please delete "methoxy" and insert therefor --methylether--.

Claim 11, line 4, please delete "methoxy" and insert therefor --methylether--.

Claim 14, line 2, delete "in that" and insert therefor --characterized in that--.

Claim 17, line 2, please delete "isocyanate";

line 3, delete "methacryloyl" and insert therefor --methacryloyl isocyanate--.

Claim 32, line 2, please delete "Claims 29 to 31" and insert therefor --Claim 29--.

Kindly insert the following claims:

--34. Solid polymer electrolyte according to Claim 2, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile and of at least one acrylic or methacrylic comonomer.

35. Process for the preparation of a methacrylonitrile polymer according to Claim 30, characterized in that it comprises a crosslinking stage after the polymerization stages.

36. Process for the preparation of a methacrylonitrile polymer according to Claim 31, characterized in that it comprises a crosslinking stage after the polymerization stages.--

IN THE ABSTRACT

Please add the Abstract of the Disclosure that is provided on a separate sheet.

REMARKS

By the present Preliminary Amendment, Applicants have amended the claims to remove the multiple dependencies in the claims. As well, the specification has been amended to correct numerous typographical error and to insert appropriate headings in conformance with practice in the United States and to add the Abstract of the Disclosure.

A favorable consideration on the merits is believed to be next in order, and is earnestly solicited.

Respectfully submitted,

BURNS, DOANE, SWECKER & MATHIS, L.L.P.

By: E. Joseph Gess
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Date: March 10, 2000

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The technical field of the invention is that of solid polymer electrolytes.

10 They can also be used for the production of the
composite electrodes of a generator.

The use of solid polymer electrolytes for
25 lithium batteries has been proposed for a long time.
This is because such electrolytes make it possible to
produce, with simple preparation processes, thin
batteries of varied shapes. The known electrolytes are
composed either of a salt dissolved in a dry polymer or
30 of a polymer swollen or gelled by a solution of a salt
in a mixture of polar aprotic organic solvents.

The known polymer matrixes can be produced based on thermoplastic materials, such as PVC (poly(vinyl chloride)), PAN (polyacrylonitrile) or PVDF (poly(vinylidene fluoride)). In the absence of solvents (or plasticizers), the mixtures between these thermoplastics and the salts form insulators or very poor ionic conductors.

After addition of solvents, they become sufficiently conducting at ambient temperatures or at temperatures below ambient temperature.

However, PVC is unstable in basic medium and cannot therefore be used in combination with lithium metal or with a lithium/ion system.

Likewise, the fluorine atoms of PVDF are reactive with respect to metallic lithium. Furthermore, it should be noted that the $\text{CH}_2\text{-CF}_2$ repeat unit of PVDF has no particular affinity with lithium salts (weak solvating nature for the Li^+ cation), which compromises its electrochemical performance.

From the viewpoint of the preparation processes, solvents with low boiling points ($< 70^\circ\text{C}$) which make possible the preparation of stable solutions are not known for PVDF, which complicates the industrial processing.

PVCs and PVDFs are disclosed in particular by Patent EP 803,925.

PAN, which corresponds to the general formula $-\text{[CH}_2\text{-CH(CN)]}_n$, is an industrial polymer used as textile fiber or as precursor for carbon fibers. It comprises a nitrile group (CN) which provides good interaction with the lithium cation (Li^+). This polymer is disclosed in particular by Patents WO 95/02314 and US 5,219,679 and partially by EP 803,925. However, this polymer has disadvantages.

Thus, when it is used in combination with a metallic lithium electrode, the interfacial resistance between lithium and electrolyte steadily increases with contact time.

This disadvantage is partly related to the presence, in the repeat unit, of a hydrogen carried by the tertiary carbon which is situated in the α position with respect to the nitrile group (CN). The highly electron-withdrawing effect of this nitrile group renders this hydrogen markedly more acidic. The result of this is a modification over time in the performance of the generator. The interfacial resistance gradually

increases, which affects the performance of the battery, in particular the power delivered. Furthermore, thermal degradation of the PAN is reflected by the release of hydrocyanic acid (HCN), which is harmful from the viewpoint of safety of use, in particular for use by the general public.

Finally, from the process viewpoint, PAN is not soluble in solvents with low boiling points. The most commonly used solvent for PAN is DMF or dimethylformamide, the boiling point of which is 152°C. Such a characteristic excludes certain processes for shaping the polymer electrolyte.

This is because one of the conventional routes for the preparation of polymer electrolytes for lithium batteries consists in dissolving a polymer, a salt and a mixture of constituent solvents of the polymer electrolyte in a common solvent, preferably with a low boiling point, and then, after having cast the mixture in the form of a film, in driving off the common solvent at atmospheric pressure or under reduced pressure without removing the constituent solvents of the plasticized polymer electrolyte. The use of a common solvent such as DMF excludes a preparation process incorporating solvents with boiling points lower than that of DMF. In other processes, the PAN + salt mixture is dissolved under warm conditions in a mixture of cyclic carbonates, before being cast in order to produce a polymer electrolyte membrane. Dissolution is carried out at an excessively high temperature which excludes, for example, the use of noncyclic carbonates.

The aim of the invention is to provide a solid polymer electrolyte which does not exhibit such disadvantages. A polymer electrolyte according to the invention is less reactive with respect to lithium than PVDF. It is more stable from the electrochemical viewpoint than PAN and less toxic than the latter from the viewpoint of the emission of hydrocyanic acid (HCN) following thermal degradation. Furthermore, the

preparation of the polymer electrolyte according to the invention is simpler because, unlike PVDF and PAN, the polymer which it employs is soluble in many conventional solvents with low boiling points, such as acetone.

The solid polymer electrolyte according to the invention comprises a polymer and a salt, which may or may not be attached to the polymer, and it is characterized in that the polymer is a methacrylonitrile polymer in the form:

- of a linear homopolymer of high mass, which may or may not be reinforced, or
- of a homopolymer, which may or may not be reinforced, rendered three-dimensional by crosslinking, or
- of a linear copolymer of high mass or of a copolymer rendered three-dimensional by crosslinking, by virtue in particular of the incorporation of at least one comonomer which makes crosslinking possible.

An electrolyte of the invention can optionally comprise, in addition, at least one aprotic liquid solvent (which will generally have a plasticizing effect) and optionally a filler.

In the continuation of the text, a methacrylonitrile polymer as defined above generally is denoted by "MANP", polymethacrylonitrile (homopolymer) is denoted by "MANHP", methacrylonitrile is denoted by "MAN", a copolymer of MAN and of a comonomer is denoted by "MANCOP", propylene carbonate is denoted by "PC" and ethylene carbonate is denoted by "EC".

Methacrylonitrile is commercially available.

Patent EP 803,925 cites the polymer MANHP but as a simple alternative to PVC and to PVDF, and from about fifteen other polymers, to be used as separator in the presence of liquid solutions of salts in organic solvents.

However, this document does not cite any example of the preparation of MANP and it is silent on

the advantages which an MANP can exhibit with respect to other polymers.

In fact, it is not possible to make direct use of any methacrylonitrile polymer in preparing a solid polymer electrolyte. This is because the solvents generally used to prepare the electrolytes (such as propylene or ethylene carbonate, which make it possible to increase the conductivity and incorporate the lithium salt) dissolve the MANP and have a strong detrimental effect on its mechanical properties.

The MANP can be a linear homopolymer MANHP having a molar mass of greater than 8×10^5 g/mol, preferably of greater than or equal to 10^6 g/mol, optionally strengthened by a filler, for example silica.

The MANP can be a homopolymer MANHP which has been completely or partially crosslinked, for example by exposure to ultraviolet radiation or by electron bombardment.

When the MANP is a homopolymer, it can comprise from 5% to 20% by mass of silica as filler.

The MANP can be a copolymer MANCOP of methacrylonitrile and of a comonomer, the said copolymer being a linear copolymer having a high molar mass or a crosslinked three-dimensional copolymer of MAN and of a comonomer which makes possible crosslinking.

The use of an MANP as defined above makes it possible to obtain a polymer electrolyte exhibiting the desired qualities (nontoxicity, improved electrochemical stability, ease of industrial manufacture) while having mechanical characteristics appropriate for a practical use. It is thus possible to produce polymer electrolyte films with a thickness of a few tens of micrometers which incorporate metal salts (for example, lithium salts) and which can be wound and/or can adopt various shapes.

As regards the toxicity, the degradation of PAN was studied by I. Ouillon et al. (Macromol. Chem.

Phys., 198, 3425-39, 1997). These authors show that the thermal degradation of PAN is reflected essentially by losses of ammonia and of hydrocyanic acid and cyclization reactions. They also studied MANHP and found that the decomposition of the latter is reflected essentially by depolymerization reactions which release the much less dangerous methacrylonitrile monomer.

In the case of the copolymers MANCOP, the choice is advantageously made of a copolymer of MAN and of at least one comonomer which makes possible the use of solvents with low boiling points while reducing the solubility in propylene carbonate or ethylene carbonate and/or of at least one comonomer which makes possible internal plasticization of the polymer by decreasing its glass transition temperature and/or of at least one comonomer which makes possible the introduction of an ionic functional group in order to obtain a unipolar electrolyte.

Such choices make it possible to obtain an electrolyte:

- the industrial use of which is rendered easier, the comonomer making possible the use of solvents with low boiling points;
- the flexibility of which is improved, the comonomer making possible internal plasticization,
- the electrochemical behavior of which is better because of the improvement in the polymer solvent interaction when the comonomer has an ionic functional group.

Mention may in particular be made, among the preferred copolymers MANCOP, of copolymers with a random tendency prepared by the radical route. They comprise one or more acrylic or methacrylic monomers. The choice of these polymers is due to reasons of copolymerization reaction kinetics, the reactivity ratios of MAN to acrylates or methacrylates being favorable to the preparation of the copolymers.

It is thus possible to use acrylic or methacrylic comonomers which make possible internal

plasticization of the polymer, the effect of which is to decrease the glass transition temperature (T_g) of the copolymer with respect to that of the homopolymer MANHP (thus to increase the flexibility of the polymer). This is because the semi-empirical Fox relationship makes it possible for a person skilled in art to predict the T_g of a copolymer in which two monomer units A and B are combined randomly according to $1/T_{gAB} = W_A/T_{gA} + W_B/T_{gB}$, in which relationship T_{gA} and T_{gB} respectively represent the glass transition temperatures of the homopolymers poly(A) and poly(B) while W_A and W_B represent the fractions by mass of the two monomer units incorporated in the copolymer.

Thus, if the comonomer A added to methacrylonitrile (monomer B) is butyl acrylate, the T_{gA} of the homopolymer poly(butyl acrylate) is 219 kelvins (-54°C). As it is known that the T_{gB} of MANHP is 393 Kelvin (120°C), the glass transition of a copolymer of these two monomers in which W_A would represent 0.2 and W_B 0.8 can be predicted by calculating the expression: $1/T_{gAB} = 0.2/219 + 0.8/393$. A value for T_{gAB} of 339 K or 66°C should therefore be found. The incorporation of butyl acrylate therefore makes possible internal plasticization, even if the result in practice is different from the theoretical prediction. For this reason, recourse to such a plasticizing comonomer makes it possible to decrease the amount of solvents added and thus to simplify the process for preparing the electrolyte and to decrease its cost, one of the functions of the added solvents being to cause so-called external plasticization in order to lower the glass transition temperature of the polymer electrolyte.

Use may be made of acrylic or methacrylic comonomers not only to produce internal plasticization (as described hereinabove) but additionally to improve the solvating properties and/or the interaction with the constituent solvents and the constituent salt of the polymer electrolyte. They are then, for example,

methacrylic macromonomers corresponding to the formula
CHX=CZ-CO-V-Y in which X = H, Z = CH₃, V = O and
Y = [(CH₂)_m-O]_pR', with m = 2, R' = H or CH₃ and
1 ≤ p ≤ 22. The poly(ethylene glycol) methacrylate sold
5 by the company Aldrich under the reference 40,953,7 can
in particular be chosen.

Use may also be made of acrylic or methacrylic
comonomers with the aim of allowing subsequent
crosslinking of the copolymer by polycondensation.
10 These monomers can have an alcohol functional group
which can be crosslinked by diisocyanate compounds,
such as toluene diisocyanate (hereinafter denoted by
"TDI") or hexamethyl diisocyanate (hereinafter denoted
by "HMDI"). One alternative consists in introducing an
15 isocyanate functional group into the copolymer by using
methacryloyl isocyanate as comonomer. The isocyanate
functional group incorporated can be condensed with
alcohol functional groups to give an urethane or with
an amine to give an urea functional group. For this
20 reason, the copolymer comprising an isocyanate
functional group can be crosslinked by polycondensation
with a polyol (crosslinking via a urethane bond),
generally a diol or a triol, or a polyamine
(crosslinking via a urea bond). A diamine or a triamine
25 will be used, for example. If the comonomer chosen is a
methacrylate comprising a trialkoxysilyl functional
group, such as (trimethoxysilyl)propyl methacrylate,
the crosslinking can be carried out subsequently by
hydrolysis/condensation of the methoxysilyl functional
30 groups in the presence or absence of other
polyalkoxysilanes. This process is conventional and is
known under the term of sol-gel process. The
crosslinking is carried out here via a siloxane bond
which is more flexible than the urethane or urea bonds.
35 The term "polyalkoxy" is understood to mean di-, tri-
or tetraalkoxy.

Use may be made of acrylic or methacrylic
comonomers which make possible subsequent crosslinking
of the material by cationic polymerization. Such

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comonomers have an oxirane functional group which does not polymerize by the radical route, as in glycidyl acrylate and methacrylate. After formation of the copolymer, the product can be photocrosslinked by cationic photoinitiation.

Finally, it is possible to provide for at least one comonomer which makes possible the introduction of an ionic functional group in order to obtain a unipolar electrolyte. Such a characteristic makes it possible to have an immobilized anion (or cation) as it is bonded to the polymer, the conductivity then being provided only by the other ion of the electrolyte.

In a specific embodiment, the polymer MANP is a polymer of MAN and of at least one acrylic comonomer corresponding to the formula $\text{CHX}=\text{CZ}-\text{CO}-\text{V}-\text{Y}$ in which:

$\text{X} = \text{C}_n\text{H}_{2n+1}$, $0 \leq n \leq 8$, preferably $n = 0$;

$\text{Z} = \text{C}_n\text{H}_{2n+1}$, $0 \leq n \leq 8$ (preferably $n = 0$ or 1), or $\text{Z} = (\text{CH}_2)_m\text{CN}$, with $0 \leq m \leq 4$,

$\text{V} = \text{O}$, NH or NR with $\text{R} = \text{C}_n\text{H}_{2n+1}$, $0 \leq n \leq 8$ (preferably $0 \leq n \leq 4$);

$\text{Y} = \text{C}_n\text{H}_{2n+1}$, with $0 \leq n \leq 8$, $\text{C}_n\text{H}_{2n}-(\text{CH}-\text{CH}_2)-\text{O}$ (oxirane functional group with $0 \leq n \leq 4$) or $[(\text{CH}_2)_m-\text{O}]_p\text{R}'$ with $m = 2, 3$ or 4 (preferably $m = 2$), $0 \leq p \leq 50$ (preferably between 2 and 22) and $\text{R}' = \text{C}_n\text{H}_{2n+1}$ with $0 \leq n \leq 8$ (preferably $n = 0, 1$ or 2).

According to the choice of the various parameters, it is possible to prepare a copolymer of MAN and of at least one alkylenebisacrylamide comonomer ($\text{V} = \text{NH}$, $\text{X} = \text{H}$, $\text{Z} = \text{H}$, $\text{Y} = (\text{CH}_2)_n-\text{NH}-\text{CO}-\text{CH}=\text{CH}_2$ ($n = 1$ to 6)), such as methylenebisacrylamide or hexamethylenebisacrylamide, or of at least one alkyl diacrylate or dimethacrylate comonomer or poly(ethylene glycol) diacrylate or dimethacrylate comonomer.

It is also possible to prepare a copolymer of MAN and of at least one poly(ethylene glycol) methoxy methacrylate (PEGMM) comonomer (i.e. $\text{V} = \text{O}$, $\text{X} = \text{H}$, $\text{Z} = \text{CH}_3$ and $\text{Y} = [(\text{CH}_2)_m-\text{O}]_p\text{R}'$ with $m = 2$ and $\text{R}' = \text{CH}_3$).

It is also possible to prepare a copolymer of MAN and of at least one hydroxyalkyl acrylate comonomer

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(i.e., $X = H$, $Z = H$, $V = O$ and $Y = [(CH_2)_m-O-]_pR'$ with $p = 1$, $R' = H$ and $1 \leq m \leq 8$ or hydroxyalkyl methacrylate comonomer (i.e., $X = H$, $Z = CH_3$, $V = O$ and $Y = [(CH_2)_m-O-]_pR'$ with $p = 1$, $R' = H$ and $1 \leq m \leq 8$).

5 The comonomer can be chosen more particularly from:

- hydroxyethyl acrylate and hydroxyethyl methacrylate ($m = 2$),
- hydroxypropyl acrylate and hydroxypropyl methacrylate ($m = 3$),
- 10 - hydroxybutyl acrylate and hydroxybutyl methacrylate ($m = 4$).

 An electrolyte can advantageously be prepared comprising at least one terpolymer of methacrylonitrile
15 (MAN), of poly(ethylene glycol) methoxy methacrylate (hereinafter denoted by "PEGMM") and of hydroxyethyl methacrylate (hereinafter denoted by "HEMA") in respective molar proportions x , y and z such that $40\% < x < 97\%$, $1\% < y < 40\%$ and $1\% < z < 20\%$, preferably $85\% < x < 90\%$,
20 $1\% < y < 7\%$ and $2\% < z < 8\%$.

 It is also possible to prepare a solid polymer electrolyte comprising at least one terpolymer of MAN, of octyl methacrylate and of HEMA in the respective molar proportions x , y and z such that $60\% < x < 97\%$,
25 $1\% < y < 30\%$ and $1\% < z < 15\%$, preferably $80\% < x < 95\%$, $1\% < y < 10\%$ and $2\% < z < 10\%$.

 It is also possible to prepare a solid polymer electrolyte comprising at least one terpolymer of MAN, of butyl methacrylate and of HEMA in the respective molar proportions x , y and z such that $50\% < x < 97\%$,
30 $1\% < y < 40\%$ and $1\% < z < 15\%$ and preferably $70\% < x < 95\%$, $1\% < y < 20\%$ and $2\% < z < 10\%$.

 Finally, it is possible to prepare an electrolyte comprising at least one terpolymer of MAN,
35 of hexyl acrylate ($X = H$, $Z = H$, $V = O$, $Y = C_6H_{13}$) and of HEMA in the respective molar proportions x , y and z such that $60\% < x < 97\%$, $1\% < y < 30\%$ and $1\% < z < 10\%$, preferably $70\% < x < 95\%$, $1\% < y < 25\%$ and $2\% < z < 8\%$.

Use may be made, as an alternative to acrylic comonomers, of vinyl monomers which are copolymerized with MAN and which are chosen, for example, so as to introduce an ionic functional group into the polymer chain. Thus, the anion is attached via a covalent bond to the polymer chain and a polymer electrolyte with cationic unipolar transport is obtained. Use may be made of a lithium styrenesulfonate, a lithium vinylsulfonate, a lithium acrylamidopropanesulfonate or a lithium N,N-diallyl-1-amidotetrafluoroethanesulfonate. The latter salt, which carries a perfluorosulfonate functional group, is disclosed in particular by Patent FR 2,687,671.

The solid polymer electrolyte can also comprise at least one bipolymer of MAN and of a monomer carrying an ionic functional group of carboxylate, phosphate, phosphonate, sulfonate or perfluorosulfonate type. More specifically, this electrolyte comprises at least one bipolymer of MAN and of a monomer carrying a perfluorosulfonate functional group chosen from lithium N,N-diallyl-1-amidotetrafluoroethanesulfonate ($\text{CH}_2=\text{CH}-\text{CH}_2)_2\text{NCOCF}(\text{CF}_3)\text{SO}_3\text{Li}$ and lithium allyloxytetrafluoroethylsulfonate $\text{CH}_2=\text{CH}-\text{CH}_2-\text{O}-(\text{CF}_2)_2-\text{SO}_3\text{Li}$.

According to another embodiment of the invention, the solid polymer electrolyte comprises a copolymer of methacrylonitrile and of at least one comonomer comprising an oxirane functional group, such as glycidyl acrylate or glycidyl methacrylate.

According to another embodiment, the solid polymer electrolyte comprises a copolymer of methacrylonitrile and of at least one acryloyl isocyanate or methacryloyl isocyanate comonomer. The molar proportions of comonomer with respect to the MAN are then preferably between 1 and 20%, more particularly between 3 and 10%.

According to another embodiment, the solid polymer electrolyte according to the invention comprises a copolymer of methacrylonitrile and of at least one (trimethoxysilyl)propyl methacrylate

comonomer in molar proportions with respect to the MAN preferably of between 1 and 40%, more particularly between 3 and 15%.

According to another embodiment of the invention, the solid polymer electrolyte according to the invention comprises a copolymer of methacrylonitrile and of at least one methacrylate comonomer carrying a cyclic carbonate functional group. This comonomer is preferably used in molar proportions with respect to the MAN of between 1 and 40%, more particularly between 5 and 25%.

According to another embodiment of the invention, the solid polymer electrolyte comprises a copolymer of methacrylonitrile and of at least one comonomer of aromatic olefin type, such as indene, acenaphthylene, vinylnaphthalene, vinylferrocene, vinylpyridine and styrene, which may or may not be substituted. This comonomer is preferably used in molar proportions with respect to the MAN of between 1 and 30%, more particularly between 5 and 15%.

According to another embodiment, the solid polymer electrolyte comprises a copolymer of methacrylonitrile and of at least one comonomer of styrene type substituted on the aromatic ring at the meta or para position with respect to the double bond by substituents R representing C_nH_{2n+1} , with $1 \leq n \leq 8$, CH_3O , CH_2Cl , CH_2Br or $C_nH_{2n+1}-OH$ with $1 \leq n \leq 8$. The molar proportions of the comonomer with respect to the MAN are preferably between 1 and 30%, more particularly between 5 and 20%.

According to another embodiment of the invention, the solid polymer electrolyte comprises a copolymer of methacrylonitrile and of at least one acrylonitrile, cyanoacrylate or cyanomethacrylate comonomer.

According to other embodiments, a polymer electrolyte can be prepared comprising a copolymer of methacrylonitrile and of at least one comonomer chosen from 1,1-dicyanoethene, vinylimidazole, vinyltriazole,

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vinyltetrazole, vinyl acetate, vinyl chloride, vinyl fluoride, vinylidene chloride, vinylidene fluoride, vinylene carbonate and maleic anhydride.

5 Mention may be made, among preferred
6 copolymers, of so-called "block" MAN copolymers.
7 Conventionally, in block copolymers, either the
8 combined physicochemical properties of the various
9 blocks (when the blocks are immiscible with one
10 another) or intermediate properties are obtained.
11 Although some block copolymerizations can be carried
12 out by the cationic route and, possibly, by the radical
13 route, anionic polymerization remains unquestionably
14 the synthetic route which is the best suited to the
15 preparation of block copolymers. However, it requires
16 so-called "living" polymerizations (quantitative and
17 rapid initiation before propagation, no transfer or
18 termination reactions). PVC and PVDF cannot be prepared
19 by the anionic route because the halogens react with
20 the very strong bases used in anionic polymerization.
21 PAN can be prepared by the anionic route but its
22 polymerization is not living. On the other hand, the
23 anionic polymerization of MAN is a living
24 polymerization. It is known that the anionic
25 polymerization of ethylene oxide is a living
26 polymerization. MANHP blocks are therefore
27 advantageously combined with more flexible
28 poly(oxyethylene) (POE) blocks, which contribute
29 flexibility to the material and the ability to solvate
30 lithium salts. Whereas the order of addition of the
31 various blocks must be carried out in anionic
32 polymerization in the direction of the increasing
33 electron affinities of the two monomers, copolymers
34 comprising MANHP-POE blocks can surprisingly be
35 prepared in both directions of addition.

36 Thus, the solid polymer electrolyte according
37 to the invention can comprise at least one diblock or
38 triblock copolymer which combines at least one
39 polymethacrylonitrile sequence with at least one
40 polyether sequence. More particularly, the solid

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polymer electrolyte according to the invention can comprise at least one poly(oxyethylene)-block-polymethacrylonitrile (POE-b-MANHP) diblock copolymer.

According to another embodiment, the polymer electrolyte according to the invention comprises at least one poly(oxyethylene)-block-polymethacrylonitrile-block-poly(oxyethylene) triblock copolymer.

According to another embodiment, the solid polymer electrolyte according to the invention comprises at least one polymethacrylonitrile-block-polyether-block-polymethacrylonitrile triblock copolymer in which the central polyether sequence is a poly(oxyethylene), a poly(oxypropylene), a poly(oxytrimethylene), a poly(oxytetramethylene), a poly(oxyethylene-oxymethylene) or an α,ω -diamino polyether.

To prepare a polymer electrolyte according to the invention, at least one solvent and at least one salt are incorporated in one of the polymers MANP described above. The solvent is chosen from propylene carbonate (PC), ethylene carbonate (EC), γ -butyrolactone, dimethoxyethane or a dialkyl carbonate. The salt is a metal salt, in particular a lithium salt chosen from lithium halides LiX ($\text{X} = \text{Cl}, \text{Br}, \text{I}$ or I_3), lithium perfluorosulfonate ($\text{C}_n\text{F}_{2n}\text{SO}_3\text{Li}$), lithium (trifluoromethylsulfonyl)imide ($\text{N}(\text{CF}_3\text{SO}_2)_2\text{Li}$), lithium bis(trifluoromethylsulfonyl)methide ($\text{HC}(\text{CF}_3\text{SO}_2)_2\text{Li}$), lithium tris(trifluoromethylsulfonyl)methide ($\text{C}(\text{CF}_3\text{SO}_2)_3\text{Li}$), lithium perchlorate (LiClO_4), lithium hexafluoroarsenate (LiAsF_6), lithium hexafluorophosphate (LiPF_6), lithium hexafluoroantimonate (LiSbF_6) and lithium tetrafluoroborate (LiBF_4).

Another subject-matter of the invention is the processes for the preparation of such polymer electrolytes.

In one embodiment, an electrolyte according to the invention is prepared by dissolving the MANP and the salt in a solvent with a low boiling point (for example, in acetone), by casting the solution onto an

appropriate substrate and by then removing the solvent. In addition, it is possible to introduce, into the solution, one or more additional solvents having a higher boiling point which act as plasticizers (for example, EC or PC); and/or a filler, such as silica, of use in particular when the MANP is a non-crosslinked homopolymer; and a crosslinking agent, when the comonomer of the MANP carries groups capable of crosslinking. When the polymer is a MANCOP obtained from a comonomer carrying ionic groups, it is not essential to dissolve a salt in the solvent.

In another embodiment, an electrolyte is prepared analogously but using only solvents, such as EC or PC, which act as plasticizers.

It is also possible to prepare an electrolyte by a process which consists in dissolving the polymer and optionally a crosslinking agent and/or fillers in acetone, in then casting the solution obtained onto an appropriate substrate, in removing the acetone, in crosslinking the film and in then incorporating therein a solution of the salt in a solvent.

In another embodiment, a dry mixture of the various constituents of the electrolyte, namely the MANP, the salt and/or the plasticizing solvents and/or a crosslinking agent, is prepared, the mixture is then extruded, for example through a flat die, and the film obtained is crosslinked.

In another alternative form, the electrolyte can be obtained in the form of a microporous membrane, for example by a phase inversion process. The MANP is dissolved in a mixture composed of a good solvent (for example, acetone) and of a non-solvent compound (for example, hexane), the non-solvent having a higher boiling point than that of the solvent. The solution obtained is subsequently cast onto an appropriate substrate and the solvent having the lowest boiling point is evaporated. After completely removing this solvent, the MANP precipitates from the non-solvent compound and forms a microporous membrane. In another

embodiment, a microporous membrane can be obtained from a solution of MANP in a solvent (for example, acetone or THF) which is cast onto an appropriate substrate, the film on the substrate thus obtained subsequently
5 being immersed in a non-solvent compound (for example, hexane) in order to bring about the precipitation of the MANP in the form a microporous membrane. In both cases, the microporous membrane can subsequently be impregnated with a liquid electrolyte comprising the
10 salt and/or solvents and/or a filler and/or a crosslinking agent. This liquid electrolyte fills the micropores of the membrane and can partially swell the MANP.

An electrolyte of the present invention can be
15 used as electrolyte in a lithium electrochemical generator, in a supercapacitor or in an electrochromic system.

A process for the bulk preparation of a methacrylonitrile homopolymer by the radical route
20 comprises the following stages:

- a thermal-decomposition free-radical initiator is dissolved in MAN,
- the mixture is degassed in order to remove the oxygen and is introduced into a hermetically closed
25 chamber,
- the mixture is brought to a temperature of 60 to 90°C and is maintained at this temperature for 24 to 72 hours.

The initiator can advantageously be chosen from
30 azobisisobutyronitrile (AIBN), benzoyl peroxide and potassium persulfate.

In a process for the preparation of a methacrylonitrile block copolymer, the polymerization of the polymethacrylonitrile block or blocks can be
35 carried out by monofunctional, difunctional or tri-functional anionic initiation from a poly(oxyethylene) block terminated by one, two or three alkoxide groups.

According to another embodiment, the poly(oxyethylene) block or blocks are obtained by mono-

or difunctional anionic initiation of the polymerization of ethylene oxide from a living anionic polymethacrylonitrile chain.

5 The processes for the preparation of a methacrylonitrile homopolymer or copolymer can comprise a crosslinking stage after the polymerization stages. The crosslinking can be carried out by irradiation or by electron bombardment, after incorporation of a photoinitiator and optionally of a monomer with a
10 functionality of 2 or 4 in a solution of MANHP in a solvent. Alternatively, the crosslinking can be carried out via urethane or siloxane bonds.

15 It is also possible to obtain a crosslinked or non-crosslinked methacrylonitrile copolymer by a process essentially comprising a stage of photochemical initiation of radical polymerization by irradiation (in particular UV irradiation). Such a process results directly in the production of a thin film of polymer electrolyte.

20 Alternatively, a methacrylonitrile homopolymer or copolymer can be prepared by radical polymerization according to conventional industrial emulsion or suspension processes.

25 The invention will be better understood on reading the description which will follow of various examples of the preparation of polymer electrolytes.

Example 1

30 6 ml of MAN (methacrylonitrile), freshly distilled under argon in order to remove oxygen, are introduced into a tube. 3 mg of initiator AIBN (azobisisobutyronitrile) are then dissolved in the MAN, without addition of solvent. The mixture is subsequently degassed in order to remove oxygen and the
35 tube is then hermetically closed and brought to 70°C for 48 hours.

The following experimental conditions were chosen in order to obtain high molar masses: moderate temperature, purification of the monomer, low amounts

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of initiator and short reaction time. In this bulk polymerization (that is to say, without solvent other than the monomer MAN), the homopolymer MANHP obtained is insoluble in the monomer MAN and forms a transparent and hard gel. After dissolution in acetone, the reaction mixture is precipitated from pentane.

After washing the precipitate with pentane and drying, a precipitated polymer is obtained with a yield of 30%. The intrinsic viscosity of the polymer was determined in the solvent DMF (dimethylformamide) at 25°C by using a capillary viscometer with an Ubbelohde tube.

A comparative study carried out between the MANHP thus synthesized and commercial MANHP supplied by the company Aldrich showed that the latter was markedly less viscous than the synthesized polymer. The Mark-Houwink-Sakurada law for MANHP in DMF has made it possible to calculate a viscometric mass $M_v = 1.1 \times 10^6$ g/mol, whereas that of the commercial MANHP is 0.45×10^6 g/mol.

By using the optimum electrolyte compositions (CP/EC/LiClO₄) given in the literature, two films were cast with the synthesized MANHP and the commercial MANHP respectively. The film prepared with the commercial MANHP is a viscous liquid which flows, whereas that prepared from the MANHP synthesized according to the above method forms a film which does not flow.

The mechanical properties of the film prepared with the synthesized MANHP can be improved by UV irradiation and/or by the addition of fillers or fibrous reinforcements. It is possible in particular to add, in proportions of 5 to 20% by mass, silica supplied by the company Degussa under the reference FK325. Such an addition makes it possible to improve the mechanical properties of the polymer electrolyte without harming its electrochemical properties.

Example 2

3% by mass, with respect to the MANHP, of a photoinitiator, such as Irgacure I 651 (2,2-dimethoxy-1,2-diphenylethane-1-one), are added to a solution in acetone of MANHP as prepared in accordance with Example 1.

After evaporating the acetone, the film is subjected to UV irradiation using a Philips UV lamp of HPA type with metal iodides (iron and cobalt iodides), the intense UV-A emission spectrum of which lies between 350 and 400 nanometers after an exposure time of 25 min. The sample is placed at approximately 8 cm from the UV lamp and cooled by a Peltier-effect system in order to keep its temperature below 50°C.

In fact, and despite the absence of unsaturations, the MANHP partially crosslinks. The level of extractables is then 85% but the behavior with respect to the conventional electrolyte is improved, probably because of branching reactions which increase the molar mass Mw of the non-crosslinked parts.

In an alternative form, 3% by mass of Irgacure I 651 and 50 to 80% by moles of ethylene carbonate as plasticizing solvent are incorporated in the sample of MANHP dissolved in the minimum amount of acetone. Evaporation of the acetone does not result in evaporation of the ethylene carbonate.

Example 3

A methylenebisacrylamide (hereinafter denoted by "MBA"), which is a monomer with a functionality of 4, in different proportions, and 3% by weight of Irgacure I 651 with respect to the MANHP are added to a concentrated solution of MANHP in acetone prepared in accordance with Example 1. The film is cast into a substrate.

After evaporating the acetone, the film is subjected to UV exposure with the lamp used in Example 2. The level of insolubles was measured after washing the membrane obtained in a large excess of

acetone. The difference between the weight of the membrane before washing and after washing, followed by drying, making it possible to calculate the level of insolubles. Tests were carried out for different % by mass of monomer MBA with respect to the MAN. The results obtained show that the proportion of insolubles increases with the level of MBA and passes through a maximum for a level of MBA of 30%. However, this level of insolubles already reaches 40% for a level of MBA of only 3%.

Example 4

An MAN/glycidyl methacrylate copolymer was prepared. Two syntheses were carried out with the following molar proportions of the monomers: the first with 95% of MAN and 5% of glycidyl methacrylate and the second with 80% of MAN and 20% of glycidyl methacrylate. The copolymerizations, carried out in cyclohexanone, are initiated by AIBN (3 mg of AIBN are used per 6 ml of solution). The synthesis takes place under argon pressure at 70°C for several days. The copolymers are obtained with a yield in the region of 80%. They exhibit a quasi-statistical distribution of the two comonomers in the macromolecular chains. However, the glycidyl methacrylate has a lower reactivity than that of the MAN and the proportion of the two monomer units in the copolymer is substantially different from that of the starting mixture of monomers. Thus, the copolymers obtained with the starting proportions given hereinabove have the following compositions: 99% MAN and 1% glycidyl methacrylate for the first synthesis, 96% MAN and 4% glycidyl methacrylate for the second. It is therefore necessary, in order to achieve a degree of incorporation of Z% of glycidyl units, to provide for a starting proportion of glycidyl methacrylate approximately equal to 5Z%. The crosslinking of such a copolymer can be carried out by opening with diamines or triamines. In the present example, the crosslinking

was obtained by photopolymerization under UV irradiation using diphenyliodonium hexafluoroantimonate $[(C_6H_5)_2I^+ SbF_6^-]$ as cationic photoinitiator. The latter releases, under UV, a Lewis acid which initiates the cationic polymerization of the oxirane rings. A level of insoluble of 63% was obtained for a proportion of 4% by mass of initiator. It is possible to increase this percentage by starting from a copolymer with a higher mass which is richer in glycidyl functional groups.

10

Example 5

An MAN/HEMA copolymer comprising 3 mol% of HEMA and 97% of MAN was synthesized. The percentages of the two monomers in the copolymer were determined by proton NMR using a Bruker 200 MHz Fourier transform spectrometer. The synthesis took place under bulk conditions at 70°C for 48 h using AIBN as initiator. 3 mg of initiator AIBN were used per 8 ml of mixture of monomers (3% HEMA and 97% MAN). The copolymer obtained is soluble in acetone and was reprecipitated from pentane. The copolymer is also soluble in THF (tetrahydrofuran), which allowed it to be characterized by gel permeation chromatography using a set of ultrastyrigel columns with porosities of 500, 10^3 , 10^4 and 10^5 angstroms, calibrated with a standard set of polystyrene samples. Mass molar masses M_w in the region of 10^5 g/mole, calculated as polystyrene equivalent, were obtained. Crosslinking via a urethane bond was carried out by using toluene diisocyanate (TDI) or hexamethyl diisocyanate (HMDI) as initiator. The TDI and the HMDI are used in a stoichiometric proportion with respect to the alcohol functional groups of the HEMA unit (number of OH functional groups of the HEMA equal to the number of NCO functional groups of TDI or HMDI, that is to say $NCO/OH = 1$). Dibutyltin dilaurate is used as catalyst in a proportion of 5 mol% with respect to the number of moles of NCO functional group. The copolymer, the diisocyanate and the catalyst are dissolved in concentrated solution in acetone or THF.

After degassing in order to remove air bubbles, the product is cast into a substrate, the solvent (acetone or THF) is evaporated and the reaction is allowed to continue for 48 hours at 30°C in a glove box maintained under dry argon.

Levels of insoluble in the region of 90% were obtained with both types of initiator. Surprisingly, crosslinking results in a decrease in the glass transition temperature by 30°C and 40°C respectively, according to whether TDI or HMDI is used. Crosslinking therefore exerts an internal plasticizing effect which makes it possible to obtain a more flexible membrane.

The MAN/HEMA copolymer could be crosslinked by treating the alcohol functional groups of the HEMA units with triethoxy(isocyanatopropyl)silane, sold by Aldrich under the reference 41,336,4. By using a 1/1 stoichiometry between the OH functional groups of the HEMA and the isocyanate functional groups of the silane, it is possible to condense the silane with the copolymer while using the same percentage of catalyst as with TDI or HMDI. The reaction product remains soluble in the reaction solvent and the functional groups can be hydrolyzed according to the conventional sol-gel mechanism (0.5 mol of water per mole of ethoxysilane functional group) to form a network with a siloxane bond. The hydrolysis/condensation reaction of the ethoxysilane bonds should preferably be carried out after having cast the film.

Example 6

Copolymers of MAN and of a poly(ethylene glycol) methoxy methacrylate which has a molar mass $M_n = 400$ g/mol and which is denoted by PEGMM 400 were synthesized, said copolymers comprising different percentages of PEGMM 400. The two monomers were mixed in the following molar proportions: for the first synthesis, 4% of PEGMM and 96% of MAN; for the second synthesis, 8% of PEGMM and 92% of MAN. 3 mg of AIBN were incorporated per 6 g of mixture of monomers.

Polymerization was carried out under bulk conditions at 70°C in sealed tube (following the protocol of Example 1). Copolymers comprising 6 mol% and 14 mol% of PEGMM 400 were thus obtained. These copolymers,
5 dissolved in acetone, are reprecipitated from pentane. Masses in the region of 10^5 g/mol were obtained

Example 7

An MAN/PEGMM/HEMA (hydroxyethyl methacrylate)
10 terpolymer comprising 2.5 mol% of PEGMM 400 and 5 mol% of HEMA were synthesized according to the same synthetic method of Example 6.

A terpolymer was obtained which is soluble in acetone. It exhibits a molecular mass of 40,000 g/mol.
15 It was subsequently crosslinked via a urethane bond by using hexamethylene diisocyanate (HMDI), which is a difunctional isocyanate, in the stoichiometric proportion (NCO/OH = 1) in the presence of dibutyltin dilaurate as catalyst (5 mol% with respect to the NCO
20 functional groups). A network exhibiting a high level of insolubles (90%) was obtained.

Example 8

0.1 mol of methacrylonitrile, 0.018 mol of
25 octyl methacrylate and 0.0062 mol of HEMA were mixed with 2 ml of butanone (solvent) in a sealed tube under argon. Radical polymerization took place at 70°C using AIBN as initiator. 3 mg of AIBN were incorporated in the mixture of monomers. The copolymer obtained is
30 dissolved in acetone and reprecipitated from pentane. A copolymer with a molar composition 69% MAN/12% HEMA/19% octyl methacrylate was obtained with a yield in the region of 40%. The number-average molecular mass recorded for this polymer is $M_n = 30,000$ g/mol. The
35 octyl methacrylate incorporated acts as internal plasticizer and contributes to a decrease in the glass transition temperature T_g ($T_g = 82^\circ\text{C}$) of approximately 40°C .

This copolymer was crosslinked via urethane bonds by using HMDI as initiator in the stoichiometric proportion ($\text{NCO/OH} = 1$) and dibutyltin dilaurate (5 mol% with respect to the NCO functional group as catalyst. A network having a level of insoluble of 86% was obtained.

Example 9

0.1 mol of methacrylonitrile, 0.021 mol of butyl methacrylate and 0.0037 mol of HEMA (hydroxyethyl methacrylate) were introduced into a sealed tube. Polymerization took place under bulk conditions at 70°C with AIBN as initiator. 3 mg of AIBN were incorporated.

This polymer is soluble in various solvents with low boiling points, such as acetone or tetrahydrofuran.

A copolymer with a molar composition of 52% MAN/9% HEMA/39% butyl methacrylate was also obtained (following a polymerization by the radical route analogous to that described in Example 8) with a yield in the region of 30%. The copolymer obtained exhibits a number-average molecular mass of 70,000 g/mol and a glass transition temperature of 80°C .

This copolymer was crosslinked via urethane bonds by using HMDI as initiator in the stoichiometric proportion ($\text{NCO/OH} = 1$) and dibutyltin dilaurate (5 mol% with respect to the NCO functional group as catalyst. A network having a level of insoluble of 85% was obtained.

Example 10

0.1 mol of MAN, 0.021 mol of hexyl acrylate and 0.0037 mol of HEMA were mixed. Polymerization took place under bulk conditions at 70°C in a sealed tube with AIBN as initiator. 3 mg of AIBN were incorporated. A copolymer with the molar composition 93% Man /2% HEMA/ 5% hexyl acrylate was obtained. It exhibits a number-average molecular mass of 50,000 g/mol and a

glass transition temperature of 80°C, lower by 40°C than that of the MANHP homopolymer.

This copolymer was crosslinked via urethane bonds by using HMDI as initiator in the stoichiometric proportion (NCO/OH = 1) and dibutyltin dilaurate (5 mol% with respect to the NCO functional group as catalyst. A network having a level of insoluble of 87% was obtained.

10

Example 11

0.09 mol of methacrylonitrile and 0.01 mol of styrene were used for a radical synthesis under bulk conditions in a sealed tube. A copolymer comprising 17% of styrene units was obtained. This copolymer is soluble in tetrahydrofuran and acetone. It exhibits a molecular mass of 60,000 g/mol.

Example 12

Various α,ω -dihydroxyoligo(oxyethylene) having different molar masses were used ($M_n = 2000, 3400, 4600$ and 8000). They are polyethylene glycols which are commercially available under the names PEG2000, PEG3400, PEG4600 and PEG8000.

After dissolving 0.69 g of PEG 4600 (i.e., an amount of OH of 3×10^{-4} mol) in 7.5 ml of THF (tetrahydrofuran), sodium hydride is added in the stoichiometric proportion (with respect to the concentration of end OHs). The solution adopts a brown coloring characteristic of the alkoxide formed. The methacrylonitrile is then added to the alkoxide in the proportions appropriate for obtaining the desired length of the MANHP blocks.

The reaction is continued for 24 hours and then it is deactivated with water. From 100 to 300 methacrylonitrile units were thus incorporated (according to the PEG used) with a yield in the vicinity of 100%.

The above procedure was used for the preparation of copolymers from PEG2000, PEG3400 and

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PEG8000 respectively. For the various syntheses carried out, the concentration in moles of OH/liter was kept constant (3×10^{-4} mol of OH) for 7.5 ml of THF.

The molar masses were determined by proton NMR and they were also measured by GPC (gel permeation chromatography). A comparison of the values obtained is summarized by the following table:

Starting PEG	Number of MAN units introduced	Mn (GPC) in g/mol	Mn (NMR) in g/mol
PEG2000	200	13,100	15,400
PEG2000	100	7700	7800
PEG3400	200	10,000	15,800
PEG4600	200	11,400	10,400
PEG4600	300	22,000	23,000
PEG7400	300	16,000	28,200

Good agreement is observed between the masses obtained by the two types of measurement. The copolymers obtained have a low polydispersity index. The ratio $I = M_w/M_n$ (M_w = mass-average molar mass, M_n = number-average molar mass) is low, less than 1.5. This result is compatible with "living" anionic polymerization (Yield: approximately 100% of polymerization).

Surprisingly, the incorporation of MAN units makes it possible to decrease the crystallinity of the poly(ethylene oxide) (POE) sequence, as is shown by the differential thermal analysis of the samples. Completely amorphous copolymers are thus obtained for 200 methacrylonitrile units incorporated. The copolymer thus obtained exhibits a single glass transition temperature situated between the two glass transition temperatures of the two homopolymers.

A glass transition temperature in the region of -20°C is obtained for copolymers comprising a POE sequence of 45 units and two side sequences each comprising 100 methacrylonitrile units. The different

copolymers synthesized are soluble in many solvents, such as acetone and tetrahydrofuran.

A number of the electrolytes were prepared by incorporating lithium salts in the polymers described above (Examples 1 to 12) and conductivity measurements were carried out.

These measurements are described in the examples hereinbelow.

The polymer is rendered ionically conducting by incorporation of a lithium salt. A lithium salt, such as lithium (trifluoromethanesulfonyl)imide (or LiTFSI) and the polymer are dissolved in acetone. The solution is then cast and the solvent evaporated. Conductivity measurements are carried out with an impedance meter using a cell with a blocking electrode made of stainless steel. Frequency analysis is carried out between 13 MHz and 5 Hz. The measurements are carried out under dynamic vacuum (dry polymer electrolyte) or under a dry argon excess pressure (plasticized polymer electrolytes).

Example 13

The conductivity of the MANHP in the dry state as prepared in Example 1 and in the presence of lithium (trifluoromethylsulfonyl)imide (LiTFSI) was measured over a wide range of concentrations.

This electrolyte exhibits a very low conductivity for moderate salt concentrations. The composition of the polymer electrolyte is expressed by the ratio CN/Li, which represents the ratio of the number (CN) of methacrylonitrile units to the number (Li) of moles of salt. A conductivity of 8×10^{-8} S/cm at 40°C is obtained for a ratio CN/Li = 8.

Contrary to the phenomenon observed for a poly(oxyethylene), the conductivity of the dry MANHP increases when the concentration of salt increases, that is to say when the CN/Li ratio decreases. For very high concentrations of salt, advantageous conductivities are obtained, such as 5×10^{-5} S/cm for a

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ratio CN/Li = 2. The LiTFSI salt therefore acts surprisingly as an external plasticizer.

CN/Li	Conductivity σ (S/cm) at 40°C	Conductivity σ (S/cm) at 80°C
2	8×10^{-7}	5×10^{-5}
3	10^{-8}	3×10^{-6}
6	10^{-8}	3×10^{-6}
8	8×10^{-8}	8×10^{-6}
12	10^{-8}	3×10^{-6}
16	-	2×10^{-6}
30	-	10^{-7}

5 This result, obtained in the absence of
solvent, shows a very good interaction between the
thermoplastic matrix and the salt, since the latter
makes it possible to prepare polymer electrolytes with
high concentrations of salt, which is favorable to the
10 conductivities of the plasticized polymer electrolytes.

Example 14

30% by mass of methylenebisacrylamide, which is
a monomer with a functionality of 4 (sold by the
15 company Aldrich under the reference 14,832-6), 3% by
mass of Irgacure I 651, the salt LiTFSI and ethylene
carbonate (EC), as plasticizer, are added to a solution
of MANHP in acetone.

20 After evaporating the acetone, the film
obtained is exposed to UV radiation with the lamp
described in Example 2. The propylene carbonate used
for the conductivity measurements is only added after
the crosslinking, in order to prevent it from being
evaporated by infrared heating during the crosslinking.

25 A conductivity of 10^{-3} S/cm at 20°C is obtained
for an electrolyte with the molar composition
MANHP (20%)/EC (50%)/PC (25%)/ LiTFSI (5%). "MANHP
(20%)" means that the proportion of homopolymer MANHP
present in the electrolyte is such that the electrolyte

comprises 20 mol of MAN units per 50 mol EC, 25 mol of PC and 5 mol of LiTFSI.

Furthermore, an electrolyte with a molar composition MANHP (25%)/EC (70%)/PC (00%)/ LiTFSI (5%) was prepared, which electrolyte exhibits an electrical conductivity of 10^{-3} S/cm at 20°C. The conductivity remains equal to 7×10^{-4} S/cm at 8°C and then it rapidly decreases at a lower temperature because of the crystallization of the EC.

Because they only employ one solvent (EC), which is the least volatile, the latter electrolytes are easier to produce but they are restricted to uses at temperatures of greater than 8°C.

Electrochemical stability studies have been carried out on the sample with a molar composition MANHP (20%)/EC (50%)/PC (25%)/ LiTFSI (5%). The sample exhibits, by cyclic voltammetry with regard to a stainless steel macroelectrode, good stability towards reduction up to 0 volt with respect to the electrochemical couple Li/Li⁺ and towards oxidation up to at least 4.1 V (still with respect to the couple Li/Li⁺). Monitoring the change, over time, in the interfacial resistance between the polymer electrolyte and the lithium metal electrode at 40°C shows good stability of the interface over several days. This behavior is different from that observed with a conventional electrolyte in which PAN, PC and LiClO₄ are combined, which electrolyte shows a continuous increase in the interfacial resistance with time.

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Table giving the ratios R/R_{t_0} of the interfacial resistances over time.

Time in hours	R/R_{t_0}	R/R_{t_0}
	Li/(PAN/EC/PC/LiClO ₄) [1]	Li/(MANHP/EC/PC/LiTFSI)
0	1	1
2	1.50	1.14
24	3	1.14
50	5	1.15
70	6	1
80	7	0.9
95	7.6	0.97
120	9	1
130	11	0.97

- 5 [1] test for PAN described in: G.B. Appetecchi, F. Croce and B. Scrosati, *Electrochimica Acta*, 40, 991, 1995

10 The initial interfacial resistance is R_{t_0} and the resistance at time t is recorded as R . The change in the ratio R/R_{t_0} over time shows, to within experimental uncertainty, the stability of the MANHP, whereas the interfacial resistance increases steadily for the PAN-based electrolyte and is found to be multiplied by a factor of more than 10 after 5 days.

15

Example 15

Conductivity measurements were carried out on a copolymer synthesized as described in Example 5 (MAN/HEMA). The copolymer, the salt LiTFSI, the ethylene carbonate, the propylene carbonate and the hexamethylene diisocyanate are dissolved in acetone. The solution is then cast on a substrate and then the acetone is evaporated. The solution is left under an inert atmosphere, in a glove box under argon, for 48 hours at 30°C in order to bring the crosslinking to completion.

20

25

The molar composition chosen for the conductivity measurements is as follows:

MAN/HEMA (15%)/EC (56%)/PC (23%)/ LiTFSI (5%) copolymer.

5 100 mol of such a material comprise 56 mol of EC, 23 mol of PC, 5 mol of LiTFSI, $97/100 \times 15 = 14.55$ mol of MAN units and $3/100 \times 15 = 0.45$ mol of HEMA units.

10 The polymer electrolyte prepared exhibits a conductivity of 8×10^{-4} S/cm at 20°C and 4×10^{-4} at 0°C.

Example 16

15 Conductivity measurements were carried out on a terpolymer synthesized as described in Example 10. The copolymer, the salt LiTFSI, the ethylene carbonate, the propylene carbonate and the hexamethylene diisocyanate are dissolved in acetone. The solution is then cast on a substrate and then the acetone is evaporated. The
20 solution is left for 48 hours in order to bring the crosslinking to completion. The molar composition used for the conductivity measurements is as follows:

MAN/HEMA/HA (15%)/EC (56%)/PC (23%)/ LiTFSI (5%) copolymer.

25 100 mol of such a material comprise 56 mol of EC, 24 mol of PC, 5 mol of LiTFSI, $93/100 \times 15 = 13.95$ mol of MAN units, $2/100 \times 15 = 0.3$ mol of HEMA units and 0.75 mol of CA (hexyl acrylate) units.

30 The conductivity is tested at 10^{-3} S/cm at 20°C and 5×10^{-4} S/cm at 0°C.

Example 17

35 Conductivity measurements were carried out on a copolymer synthesized as described in Example 12. This copolymer comprises a POE sequence with 45 units and two side sequences each comprising 100 methacrylonitrile units. The copolymer is dissolved in acetone in the presence of the lithium salt (LiTFSI). The film is cast and the solvent is then evaporated.

The polymer film is left to dry for several days. In this type of copolymer, the POE block and the MANHP block can contribute to solvating the Li^+ cation. The composition of the electrolyte is therefore expressed as $(\text{O}+\text{CN})/\text{Li}$, the ratio of, on the one hand, the sum of the number of moles of oxyethylene units (O) and of the number of moles of MAN (CN) to, on the other hand, the number of moles of lithium salt (Li).

The dry electrolyte obtained exhibits a conductivity of 10^{-6} S/cm at approximately 25°C for a concentration of salt $(\text{O}+\text{CN})/\text{Li} = 8$. If the concentration of salt in the electrolyte is increased to the ratio $(\text{O}+\text{CN})/\text{Li} = 1$, the conductivity increases to 2×10^{-5} S/cm at 20°C . This value is equivalent to those which are obtained with the best amorphous polyethers.

The salt LiTFSI therefore acts here as external plasticizer. This surprising phenomenon is contrary to that commonly observed for polyethers (POE), where the salt increases the T_g as a result of the strong salt-matrix interactions.

Example 18

Polymethacrylonitrile-co-poly(hydroxyethyl methacrylate) bipolymers were prepared by bulk radical copolymerization at 80°C of methacrylonitrile (MAN) and hydroxyethyl methacrylate (HEMA).

6 ml of MAN, 10 microliters of a 0.1 g/ml solution of AIBN in acetone (i.e. 1 mg) and 0.260 ml of HEMA were used. The reaction mixture was cooled after reacting for 48 hours, dissolved in acetone and then precipitated from pentane. The recovered precipitate was washed with pentane and then dried under vacuum at 50°C .

The process was repeated under the same conditions, using successively 6 ml of MAN and 0.434 ml of HEMA and then 6 ml of MAN and 0.867 ml of HEMA.

The crosslinking of the copolymers obtained makes it possible to improve their mechanical behavior

and to minimize creep problems. The copolymers were crosslinked using hexamethylene diisocyanate (HMDI). The crosslinking results from the polycondensation between the OH functional groups of the copolymer and the isocyanate functional groups of HMDI, which form urethane bonds. A solution was prepared by introducing an MAN/HEMA copolymer and HMDI into acetone in stoichiometric proportions ($\text{NCO/OH} = 1$) in the presence of dibutyltin dilaurate (5% with respect to the OH functional groups) as catalyst. After degassing in order to remove air bubbles, the solution was cast into a substrate, the solvent was evaporated and the reaction was allowed to continue for 48 hours at 30°C in a glove box maintained under dry argon.

The composition of each of the copolymers before crosslinking was determined by proton NMR. The glass transition temperature of each of the copolymers before and after crosslinking was determined. The results are collated in the table hereinbelow. "MAN/HEMA" represents the initial percentage in moles of the monomers used. "MAN/HEMA (NMR)" denotes the composition of the copolymer, as % of units, determined by NMR, "Tg(C)" denotes the glass transition temperature in degrees Celsius of the copolymers obtained and "Tg(CL)" denotes the transition temperature of the networks obtained by crosslinking the copolymers.

MAN/HEMA	MAN/HEMA (NMR)	Tg(C)	Tg(CL)
97/3	94.2/5.8	92.8	49.2
95/5	93/7	78.8	45.7
90/10	82.6/17.4	95.8	41.1

It is found that the Tg of a network is markedly less than the Tg of the corresponding copolymer.

Conductivity measurements were carried out on the three copolymers synthesized. For each of the copolymers, a solution of copolymer in acetone was

prepared and then the salt LiTFSI, the ethylene carbonate, the propylene carbonate and the HMDI were added in proportions such that the final membrane exhibits the following composition: 11% of crosslinked copolymer, 73 of EP/PC (1/1) and 16% of salt LiTFSI. The solution was cast in a glass ring with a teflon-coated base and then the acetone was evaporated. After having maintained the unit at ambient temperature for 48 hours, it was found that 90% of the copolymer had crosslinked.

The change in the conductivity as a function of the temperature was measured. The curve representing "log(conductivity)" = $f(1/\text{temperature in K})$ shows excellent behavior in a temperature range from 253 K to 363 K. The membrane undergoes neither detrimental change nor creep at high temperature, despite the high levels of solvents used. The conductivity, expressed in millisiemens/cm, reaches 0.7, 3.5 and 9 respectively at the temperatures of -20°C, +20°C and +90°C.

The membrane prepared from the 95/5 bipolymer was studied by cyclic voltammetry at 25°C using a stainless steel working electrode, a secondary electrode and a lithium reference electrode. The voltammogram obtained shows good stability in the range 0 to 4.4 volts with respect to lithium. The reoxidation/deposition of lithium efficiency is 75% under these conditions.

Example 19

Polymethacrylonitrile-co-hydroxyethyl methacrylate-co-butyl methacrylate (MAN/HEMA/BUMA) terpolymers were prepared according to the procedure described in Example 18. 6 ml of MAN, 10 microliters of a 0.1 g/l solution of AIBN in acetone, i.e. 1 mg of AIBN, 0.26 ml of HEMA and 0.569 ml of BUMA were used. The reaction mixture was cooled after reacting for 48 hours, dissolved in acetone and then precipitated from pentane. The precipitate recovered was washed with pentane and then dried under vacuum at 50°C.

The process was repeated under the same conditions using successively 6 ml of MAN, 0.26 ml and 1.137 ml of HEMA and of BUMA and then 6 ml of MAN, 0.26 ml and 2.275 ml of HEMA and of BUMA.

5 The composition of the terpolymers was determined by proton NMR on samples dissolved in deuterated D6-acetone. A solution of the terpolymers in THF was subjected to analysis by gel permeation chromatography using a set of ultrastyrigel columns
10 with porosities of 500, 10^3 , 10^4 and 10^5 angstroms. The average masses Mw and Mn were calculated in polystyrene equivalents.

It is apparent that the monomer BUMA exerts an internal plasticizing effect on the non-crosslinked
15 terpolymer, since the Tg values decrease when the level of BUMA increases.

The copolymers were crosslinked using HMDI under conditions similar to those described in Example 19. It is found that crosslinking by HMDI
20 exerts a greater plasticizing effect than incorporation of BUMA.

The results are collated in the table hereinbelow. "MAN/HEMA/BUMA" represents the initial percentage in moles of the monomers used.
25 "MAN/HEMA/BUMA (NMR)" denotes the composition of the copolymer, as % of units, determined by NMR, "Tg(C)" denotes the glass transition temperature in °C of the copolymers obtained and "Tg(CL)" denotes the transition temperature of the networks obtained by crosslinking
30 the copolymers.

MAN/HEMA/ BUMA	MAN/HEMA/BUMA [lacuna]	Mn g/mol	Mw g/mol	I=Mw/Mn	Tg (C)	Tg (CL)
92/3/5	87.2/5.5/7.3	74,800	89,000	1.2	98.4	57.7
87/3/10	78.8/8/13.2	129,900	188,900	1.45	90.8	63.5
77/3/20	72/6.4/21.6	307,900	599,300	1.9	80.4	66.7

Membranes were prepared under conditions analogous to those described in Example 18 and their

conductivity was measured. The membrane obtained from a composition comprising 73 mol% of a 1/1 EC/PC mixture, 11 mol% of network obtained from the mixture of monomers 92/3/5 and 16 mol% of $\text{LiN}(\text{SO}_2\text{CF}_3)_2$ exhibits the best conductivity (2,3 millisiemens at 20°C). A transport number for the lithium of 0.79 was determined at 25°C .

Example 20

Terpolymers in which MAN and HEMA are combined with a methacrylic monomer with a pendant oligoether chain were prepared under the same conditions as those of Example 19, the butyl methacrylate being replaced by a poly(ethylene glycol) ethyl ether methacrylate with a molar mass of 246 sold by the company Aldrich under the reference 40,954-5.

6 ml of MAN, 10 microliters of a 0.1 g/l solution of AIBN in acetone (i.e. 1 mg of AIBN), 0.17 ml of HEMA and 0.46 ml of PEGEEM were used. The reaction mixture was cooled after reacting for 48 hours, dissolved in acetone and then precipitated from pentane. The precipitate recovered was washed with pentane and then dried under vacuum at 50°C . The process was repeated under the same conditions using successively 6 ml of MAN, 0.17 ml and 0.69 ml of HEMA and of PEGEEM and then 6 ml of MAN, 0.17 ml and 0.91 ml of HEMA and of PEGEEM.

The average molar masses and the glass transition temperatures were determined. The results are collated in the table hereinbelow.

The results are collated in the table hereunder. "MAN/HEMA/PEGEEM" represents the starting percentage in moles of the monomers used. "MAN/HEMA/PEGEEM (NMR)" denotes the composition of the copolymer, as % of units, determined by NMR, "Tg(C)" denotes the glass transition temperature in $^\circ\text{C}$ of the copolymers obtained and "Tg(CL)" denotes the transition temperature of the networks obtained by crosslinking the copolymers.

MAN/HEMA/ PEGEEM	MAN/HEMA/ PEGEEM [lacuna]	Mn g/mol	Mw g/mol	I = Mw/Mn	Tg (C)	Tg (CL)
93.5/2.8/3.7	90.2/5.1/4.7	45,200	184,300	4.1	77.8	56.5
91.6/2.8/5.6	85.6/9.4/5	38,000	152,000	3.99	67.7	49.5

The conductivities at 20°C vary between 2.5 and 3.1 millisiemens/cm according to the composition of the copolymer. A transport number for the lithium of 0.57 was measured for the terpolymer with the MAN/HEMA/PEGEEM composition of 85.6/9.4/5.

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CLAIMS

1. Solid polymer electrolyte, characterized in that it comprises at least one methacrylonitrile polymer chosen from:
- linear homopolymers of high mass, which may or may not be reinforced;
 - homopolymers, which may or may not be reinforced, rendered three-dimensional by crosslinking;
 - linear methacrylonitrile copolymers of high mass;
 - copolymers of methacrylonitrile and of at least one comonomer which makes possible crosslinking, which copolymers are rendered three-dimensional by crosslinking.
2. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile and of a comonomer which makes possible the use of solvents with low boiling points and/or of a comonomer which makes possible internal plasticization of the polymer by decreasing its glass transition temperature and/or of a comonomer which makes possible the introduction of the ionic functional group in order to obtain a unipolar electrolyte.
3. Solid polymer electrolyte according to Claim 1, characterized in that it comprises a methacrylonitrile homopolymer and from 5% to 20% by mass of silica.
4. Solid polymer electrolyte according to either of Claims 1 and 2, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile and of at least one acrylic or methacrylic comonomer.
5. Solid polymer electrolyte according to Claim 4, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile and of at least one acrylic comonomer corresponding to the formula $\text{CHX}=\text{CZ}-\text{CO}-\text{V}-\text{Y}$, in which:
- X represents $\text{C}_n\text{H}_{2n+1}$, with $0 \leq n \leq 8$;

- Z represents C_nH_{2n+1} , with $0 \leq n \leq 8$, or $(CH_2)_mCN$, with $0 \leq m \leq 4$;
- V represents O, NH or NR, R represents C_nH_{2n+1} , with $0 \leq n \leq 8$;
- 5 Y represents a C_nH_{2n+1} radical, with $0 \leq n \leq 8$, a radical carrying an oxirane group $C_nH_{2n}-(CH-CH_2)-O$, with $1 \leq n \leq 4$, or a radical $[(CH_2)_m-O]_pR'$, in which $m = 2, 3$ or 4 , $1 \leq p \leq 50$ and R' represents C_nH_{2n+1} , with $0 \leq n \leq 8$.
- 10 6. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile and of an alkylenebisacrylamide corresponding to the formula $CH_2=CH-CO-NH-(CH_2)_n-NH-CO-CH=CH_2$ in which $0 \leq n \leq 6$,
- 15 7. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile and of an alkyl diacrylate or dimethacrylate or of poly(ethylene glycol) diacrylate or dimethacrylate.
- 20 8. Solid polymer electrolyte according to Claim 5, characterized in that the comonomer is a poly(ethylene glycol) methoxy methacrylate $CH_2=C(CH_3)-COO-(CH_2)_2-O-]_pCH_3$ with $1 \leq p \leq 50$.
- 25 9. Solid polymer electrolyte according to Claim 5, characterized in that the comonomer is a hydroxyalkyl acrylate corresponding to the formula $CH_2=CH-CO-O-(CH_2)_m-OH$ with $1 \leq m \leq 8$ or a hydroxyalkyl methacrylate corresponding to the formula $CH_2=C(CH_3)-CO-O-(CH_2)_m-OH$ with $1 \leq m \leq 8$.
- 30 10. Solid polymer electrolyte according to Claim 9, characterized in that the comonomer is chosen from hydroxyethyl acrylate, hydroxypropyl acrylate, hydroxybutyl acrylate, hydroxyethyl methacrylate, hydroxypropyl methacrylate and hydroxybutyl
- 35 methacrylate.
11. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a terpolymer of methacrylonitrile (MAN), of poly(ethylene glycol) methoxy methacrylate (PEGMM) and

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of hydroxyethyl methacrylate (HEMA) in the respective molar proportions x, y and z such that $40\% < x < 97\%$, $1\% < y < 40\%$ and $1\% < z < 20\%$.

12. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a terpolymer of methacrylonitrile (MAN), of octyl methacrylate or of hexyl acrylate, and of hydroxyethyl methacrylate (HEMA) in the respective molar proportions x, y and z such that $60\% < x < 97\%$, $1\% < y < 30\%$ and $1\% < z < 15\%$.

13. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a terpolymer of methacrylonitrile (MAN), of butyl methacrylate and of hydroxyethyl methacrylate (HEMA) in the respective [lacuna] proportions x, y and z such that $50\% < x < 97\%$, $1\% < y < 40\%$ and $1\% < z < 15\%$.

14. Solid polymer electrolyte according to Claim 2, in that the methacrylonitrile polymer is a bipolymer of methacrylonitrile and of a monomer carrying a carboxylate, phosphate, phosphonate, sulfonate or perfluorosulfonate ionic functional group.

15. Solid polymer electrolyte according to Claim 14, characterized in that the monomer carrying an ionic functional group is lithium N,N-diallyl-1-amidotetrafluoroethanesulfonate $(\text{CH}_2=\text{CH}-\text{CH}_2)_2\text{NCO}-\text{CF}(\text{CF}_3)\text{SO}_3\text{Li}$ or lithium allyloxytetrafluoroethyl-sulfonate $\text{CH}_2=\text{CH}-\text{CH}_2-\text{O}-(\text{CF}_2)_2-\text{SO}_3\text{Li}$.

16. Solid polymer electrolyte according to Claim 4, characterized in that the comonomer is glycidyl acrylate or glycidyl methacrylate.

17. Solid polymer electrolyte according to Claim 4, characterized in that the comonomer is an isocyanate acryloyl or methacryloyl used in molar proportions with respect to the methacrylonitrile of between 1 and 20%.

18. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile (MAN) and of at least one (trimethoxysilyl)propyl methacrylate comonomer, the molar proportion of comonomer with respect to the MAN being between 1 and 40%.

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19. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile (MAN) and of at least one methacrylate comonomer carrying a cyclic carbonate functional group, the molar proportion of comonomer with respect to the MAN being between 1 and 40%.

20. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile (MAN) and of at least one aromatic olefin chosen from the group consisting of indene, acenaphthylene, vinylnaphthalene, vinylferrocene, vinylpyridine and styrene, which may or may not be substituted, the molar proportion of comonomer with respect to the MAN being between 1 and 30%.

21. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile (MAN) and of a comonomer of styrene type substituted on the aromatic ring at the meta or para position with respect to the double bond by substituents chosen from CH_3O , CH_2Cl , CH_2Br , alkyl radicals having from 1 to 8 carbon atoms and hydroxyalkyl radicals having from 1 to 8 carbon atoms, the molar proportion of comonomer with respect to the MAN being between 1 and 30%.

22. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile (MAN) and of a comonomer of acrylonitrile, cyanoacrylate or cyanomethacrylate type.

23. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is a copolymer of methacrylonitrile (MAN) and of a vinyl comonomer chosen from the group consisting of 1,1-dicyanoethene, vinylimidazole, vinyltriazole, vinyltetrazole, vinyl acetate, vinyl chloride, vinyl fluoride, vinylidene chloride, vinylidene fluoride, vinylene carbonate and maleic anhydride.

24. Solid polymer electrolyte according to Claim 1, characterized in that the methacrylonitrile polymer is

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a diblock or triblock copolymer comprising at least one polymethacrylonitrile sequence and at least one polyether sequence.

25. Solid polymer electrolyte according to
5 Claim 24, characterized in that the copolymer is a poly(oxyethylene)-block-polymethacrylonitrile (POE-b-MANHP) diblock copolymer.

26. Solid polymer electrolyte according to
10 Claim 24, characterized in that the copolymer is a poly(oxyethylene)-block-polymethacrylonitrile-block-poly(oxyethylene) triblock copolymer.

27. Polymer electrolyte according to Claim 1,
15 characterized in that it comprises at least one solvent chosen from propylene carbonate (PC), ethylene carbonate (EC), γ -butyrolactone, dimethoxyethane and dialkyl carbonates.

28. Polymer electrolyte according to Claim 1,
20 characterized in that it comprises at least one lithium salt chosen from the group consisting of lithium halides, lithium perfluorosulfonate, lithium (trifluoromethylsulfonyl)imide, lithium bis(trifluoromethylsulfonyl)methide, lithium tris(trifluoromethylsulfonyl)methide, lithium perchlorate, lithium hexafluoroarsenate, lithium hexafluorophosphate, lithium
25 hexafluoroantimonate and lithium tetrafluoroborate.

29. Process for the bulk preparation of a methacrylonitrile polymer by the radical route, characterized in that it comprises the following stages:

30 - a thermal-decomposition free-radical initiator is dissolved in methacrylonitrile or a mixture of methacrylonitrile with at least one comonomer,

- the mixture is degassed in order to remove the oxygen and is introduced into a hermetically closed
35 chamber,

- the mixture is brought to a temperature of 60 to 90°C and is maintained at this temperature for 24 to 72 hours.

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30. Process for the preparation of a block copolymer of methacrylonitrile and of ethylene oxide, characterized in that the polymerization of the polymethacrylonitrile block or blocks is carried out by monofunctional, difunctional or trifunctional anionic initiation from a poly(oxyethylene) block terminated by one, two or three alkoxide groups.

31. Process for the preparation of a block copolymer of methacrylonitrile and of ethylene oxide, characterized in that the poly(oxyethylene) block or blocks are obtained by mono- or difunctional anionic initiation of the polymerization of ethylene oxide from a living anionic polymethacrylonitrile chain.

32. Process for the preparation of a methacrylonitrile polymer according to Claims 29 to 31, characterized in that it comprises a crosslinking stage after the polymerization stages.

33. Process for the preparation of a crosslinked or non-crosslinked methacrylonitrile polymer, characterized in that it comprises a stage of photochemical initiation of radical polymerization by UV radiation which results directly in a thin film of polymer electrolyte.

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The invention concerns a solid polymer electrolyte characterised in that it comprises at least a methacrylonitrile polymer in the form: of a linear homopolymer with strong mass, reinforced or not; or a homopolymer, reinforced or not made three-dimensional by crosslinking; or a linear copolymer with strong mass or made three-dimensional by crosslinking, in particular by incorporation of at least a crosslinkable comonomer. The invention is useful for making electrochemical generators, high load capacitors or eletrochrome systems.

COMBINED DECLARATION FOR PATENT APPLICATION AND POWER OF ATTORNEY
(Includes Reference to Provisional and PCT International Applications)ATTORNEY'S DOCKET NUMBER
032013-024

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name;
 I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

"Solid polymer electrolyte and processes for its preparation".

the specification of which (check only one item below):

☐ is attached hereto.

☒ was filed as United States application

Number 09/508,378

on March 10, 2000

and was amended

on _____ (if applicable).

☒ was filed as PCT international application

Number PCT/FR99/01680

on July 9, 1999

and was amended under PCT Article 19

on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose to the Office all information known to me to be material to patentability as defined in Title 37, Code of Federal Regulations, §1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, §119 (a)-(e) of any foreign application(s) for patent or inventor's certificate or of any PCT international application(s) designating at least one country other than the United States of America listed below and have also identified below any foreign application(s) for patent or inventor's certificate or any PCT international application(s) designating at least one country other than the United States of America filed by me on the same subject matter having a filing date before that of the application(s) of which priority is claimed:

PRIOR FOREIGN/PCT APPLICATION(S) AND ANY PRIORITY CLAIMS UNDER 35 U.S.C. §119:

COUNTRY (if PCT, indicate "PCT")	APPLICATION NUMBER	DATE OF FILING (day, month, year)	PRIORITY CLAIMED UNDER 35 U.S.C. §119
FR	98.09385	10/07/1998	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No
			<input type="checkbox"/> Yes <input type="checkbox"/> No

I hereby claim the benefit under Title 35, United States Code § 119(e) of any United States provisional application(s) listed below.

(Application Number)

(Filing Date)

(Application Number)

(Filing Date)

COMBINED DECLARATION FOR PATENT APPLICATION AND POWER OF ATTORNEY (CONTINUED)
(includes Reference to Provisional and PCT International Applications)

 ATTORNEY'S DOCKET NO.
 032013-024

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) or PCT international application(s) designating the United States of America that is/are listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in that/those prior application(s) in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose to the Office all information known to me to be material to the patentability as defined in Title 37, Code of Federal Regulations §1.56, which became available between the filing date of the prior application(s) and the national or PCT international filing date of this application:

PRIOR U.S. APPLICATIONS OR PCT INTERNATIONAL APPLICATIONS DESIGNATING THE U.S. FOR BENEFIT UNDER 35 U.S.C. 120:

U.S. APPLICATIONS		STATUS (check one)		
U.S. APPLICATION NUMBER	U.S. FILING DATE	PATENTED	PENDING	ABANDONED
PCT/FR99/01680				
PCT APPLICATIONS DESIGNATING THE U.S.				
PCT APPLICATION NO.	PCT FILING DATE	U.S. APPLICATION NUMBERS ASSIGNED (if any)		

I hereby appoint the following attorneys and agent(s) to prosecute said application and to transact all business in the Patent and Trademark Office connected therewith and to file, prosecute and to transact all business in connection with international applications directed to said invention:

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Regis E. Slutter	26,999	T. Gene Dillahunty	25,423		
Samuel C. Miller, III	27,360	Patrick C. Keane	32,858		
Ralph L. Freeland, Jr.	16,110	Bruce J. Boggs, Jr.	32,344		

and:

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

COMBINED DECLARATION FOR PATENT APPLICATION AND POWER OF ATTORNEY (CONTINUED) (Includes Reference to Provisional and PCT International Applications)		ATTORNEY'S DOCKET NO. 032013-024	
FULL NAME OF SOLE OR FIRST INVENTOR Jean-Yves SANCHEZ		SIGNATURE <i>Sanchez</i>	
RESIDENCE Le Chaboud, 781, Chemin de Chartreuse, 38330 SAINT ISMIER		DATE 27/04/00	
POST OFFICE ADDRESS Idem		CITIZENSHIP French	
FRANCE		FRX	
FULL NAME OF SECOND JOINT INVENTOR, IF ANY Fanny ALLOIN		SIGNATURE <i>Alloin</i>	
RESIDENCE 77c, avenue Jeanne d'Arc, 38100 GRENOBLE, FRANCE		DATE 27/04/00	
POST OFFICE ADDRESS Idem		CITIZENSHIP French	
FRANCE		FRX	
FULL NAME OF THIRD JOINT INVENTOR, IF ANY		SIGNATURE	
RESIDENCE		DATE	
POST OFFICE ADDRESS		CITIZENSHIP	
FULL NAME OF FOURTH JOINT INVENTOR, IF ANY		SIGNATURE	
RESIDENCE		DATE	
POST OFFICE ADDRESS		CITIZENSHIP	
FULL NAME OF FIFTH JOINT INVENTOR, IF ANY		SIGNATURE	
RESIDENCE		DATE	
POST OFFICE ADDRESS		CITIZENSHIP	
FULL NAME OF SIXTH JOINT INVENTOR, IF ANY		SIGNATURE	
RESIDENCE		DATE	
POST OFFICE ADDRESS		CITIZENSHIP	
FULL NAME OF SEVENTH JOINT INVENTOR, IF ANY		SIGNATURE	
RESIDENCE		DATE	
POST OFFICE ADDRESS		CITIZENSHIP	
FULL NAME OF EIGHTH JOINT INVENTOR, IF ANY		SIGNATURE	
RESIDENCE		DATE	
POST OFFICE ADDRESS		CITIZENSHIP	
FULL NAME OF NINTH JOINT INVENTOR, IF ANY		SIGNATURE	
RESIDENCE		DATE	
POST OFFICE ADDRESS		CITIZENSHIP	